

1000562

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* * * * * Welcome to STN International * * * * *

NEWS	1		Web Page URLs for STN Seminar Schedule - N. America
NEWS	2	Jan 25	BLAST(R) searching in REGISTRY available in STN on the Web
NEWS	3	Jan 29	FSTA has been reloaded and moves to weekly updates
NEWS	4	Feb 01	DKILIT now produced by FIZ Karlsruhe and has a new update frequency
NEWS	5	Feb 19	Access via Tymnet and SprintNet Eliminated Effective 3/31/02
NEWS	6	Mar 08	Gene Names now available in BIOSIS
NEWS	7	Mar 22	TOXLIT no longer available
NEWS	8	Mar 22	TRCTHERMO no longer available
NEWS	9	Mar 28	US Provisional Priorities searched with P in CA/CAPLUS and USPATFULL
NEWS	10	Mar 28	LIPINSKI/CALC added for property searching in REGISTRY
NEWS	11	Apr 02	PAPERCHEM no longer available on STN. Use PAPERCHEM2 instead.
NEWS	12	Apr 08	"Ask CAS" for self-help around the clock
NEWS	13	Apr 09	BEILSTEIN: Reload and Implementation of a New Subject Area
NEWS	14	Apr 09	ZDB will be removed from STN
NEWS	15	Apr 19	US Patent Applications available in IFICDB, IFIPAT, and IFIUDB
NEWS	16	Apr 22	Records from IP.com available in CAPLUS, HCAPLUS, and ZCAPLUS
NEWS	17	Apr 22	BIOSIS Gene Names now available in TOXCENTER
NEWS	18	Apr 22	Federal Research in Progress (FEDRIP) now available
NEWS EXPRESS			February 1 CURRENT WINDOWS VERSION IS V6.0d, CURRENT MACINTOSH VERSION IS V6.0a(ENG) AND V6.0Ja(JP), AND CURRENT DISCOVER FILE IS DATED 05 FEBRUARY 2002
NEWS HOURS			STN Operating Hours Plus Help Desk Availability
NEWS INTER			General Internet Information
NEWS LOGIN			Welcome Banner and News Items
NEWS PHONE			Direct Dial and Telecommunication Network Access to STN
NEWS WWW			CAS World Wide Web Site (general information)

Enter NEWS followed by the item number or name to see news on that specific topic.

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* * * * * STN Columbus * * * * *

FILE 'HOME' ENTERED AT 12:27:31 ON 09 MAY 2002

Kamal Saeed

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=> file reg

COST IN U.S. DOLLARS

SINCE FILE

ENTRY

TOTAL

SESSION

FULL ESTIMATED COST

0.21

0.21

FILE 'REGISTRY' ENTERED AT 12:27:47 ON 09 MAY 2002

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STRUCTURE FILE UPDATES: 7 MAY 2002 HIGHEST RN 412267-09-5

DICTIONARY FILE UPDATES: 7 MAY 2002 HIGHEST RN 412267-09-5

TSCA INFORMATION NOW CURRENT THROUGH July 7, 2001

Please note that search-term pricing does apply when
conducting SmartSELECT searches.

Crossover limits have been increased. See HELP CROSSOVER for details.

Calculated physical property data is now available. See HELP PROPERTIES
for more information. See STNote 27, Searching Properties in the CAS

Registry File, for complete details:

<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>Testing the current file..... screen

ENTER SCREEN EXPRESSION OR (END):end

=>

Uploading C:\Program Files\Stnexp\Queries\0989932211.str

L1 STRUCTURE UPLOADED

=> que L1

L2 QUE L1

=> s l1

SAMPLE SEARCH INITIATED 12:28:09 FILE 'REGISTRY'

SAMPLE SCREEN SEARCH COMPLETED - 6 TO ITERATE

100.0% PROCESSED 6 ITERATIONS

1 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 6 TO 266

PROJECTED ANSWERS: 1 TO 80

L3 1 SEA SSS SAM L1

=> s l1 full

FULL SEARCH INITIATED 12:28:21 FILE 'REGISTRY'

FULL SCREEN SEARCH COMPLETED - 101 TO ITERATE

100.0% PROCESSED 101 ITERATIONS

22 ANSWERS

SEARCH TIME: 00.00.01

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L4 22 SEA SSS FUL L1

=> file caplus
COST IN U.S. DOLLARS

SINCE FILE	TOTAL
ENTRY	SESSION
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FULL ESTIMATED COST

FILE 'CAPLUS' ENTERED AT 12:28:40 ON 09 MAY 2002
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FILE COVERS 1907 - 9 May 2002 VOL 136 ISS 19
FILE LAST UPDATED: 7 May 2002 (20020507/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

CAS roles have been modified effective December 16, 2001. Please check your SDI profiles to see if they need to be revised. For information on CAS roles, enter HELP ROLES at an arrow prompt or use the CAS Roles thesaurus (/RL field) in this file.

=> s l4

L5 5 L4

=> d ibib abs hitstr tot

1000562

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS
 ACCESSION NUMBER: 1994:134462 CAPLUS
 DOCUMENT NUMBER: 120:134462
 TITLE: Heterocyclic phenoxyacetic acid derivative
 antithrombotic and antihypertensive agents
 INVENTOR(S): Hamanaka, Nobuyuki; Takahashi, Kanji; Tokumoto,
 Hidekado
 PATENT ASSIGNEE(S): Ono Pharmaceutical Co., Ltd., Japan
 SOURCE: Eur. Pat. Appl., 112 pp.
 CODEN: EPXXDW
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
EP 558062	A2	19930901	EP 1993-103113	19930226
EP 558062	A3	19940112		
EP 558062	B1	19970507		

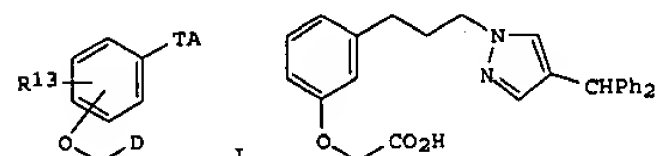
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IE, IT, LI, LU, MC, NL, PT,

SE

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AA	A2	E	T3	A	A2	JP	US	US	JP	US	US	US	JP	US	US	US	JP	US	US	
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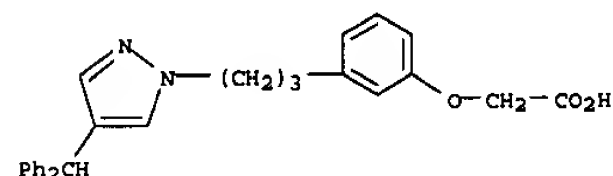
PRIORITY APPLN. INFO.: JP 1992-78330 A 19920228
 JP 1993-59418 A3 19930225
 US 1993-24306 A3 19930301
 US 1994-293218 A3 19940819
 US 1996-642598 A3 19960503

OTHER SOURCE(S): MARPAT 120:134462
 GI

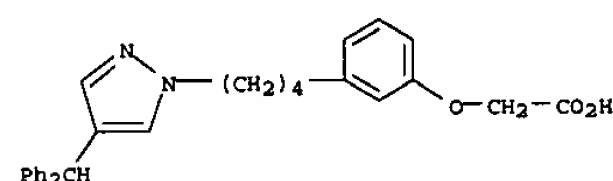


AB The title compds. I [A = heterocyclyl, carboxylate, (un)substituted CH2NH2, etc.; D = CO2R10, CONR11R12; R10 = H, C1-12 alkyl; R11, R12 = H, C1-4 alkyl; R13 = H, C1-4 alkyl, C1-4 alkoxy, NO2; T = direct bond, C1-6 alkylene, C2-6 alkenylene, O(CH2)s; s = 2-4], useful in the treatment of thrombosis, arteriosclerosis, ischemic heart disease, gastric ulcer, or hypertension, are prepd. and I-contg. formulations are presented. Thus,

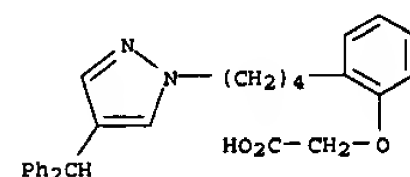
L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)
 Me 3-[3-(4-(diphenylmethyl)pyrazol-1-yl)propyl]phenoxyacetate was hydrolyzed, producing pyrazole deriv. II which demonstrated a 50% human blood platelet aggregation inhibitory concn. of 0.42 .mu.M.
 IT 152381-30-1 152381-31-2 152381-35-6
 152381-37-8 152381-38-9 152381-40-3
 152381-41-4 152381-42-5 152381-46-9
 153183-95-0 153183-96-1
 RL: RCT (Reactant)
 (antithrombotic and antihypertensive activity of)
 RN 152381-30-1 CAPLUS
 CN Acetic acid, [3-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-(9CI) (CA INDEX NAME)



RN 152381-31-2 CAPLUS
 CN Acetic acid, [3-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-(9CI) (CA INDEX NAME)

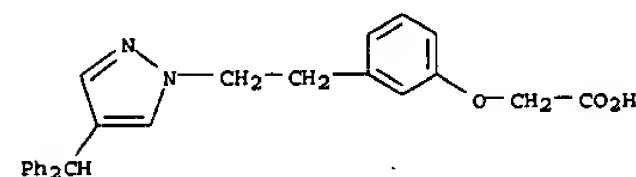


RN 152381-35-6 CAPLUS
 CN Acetic acid, [2-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-(9CI) (CA INDEX NAME)

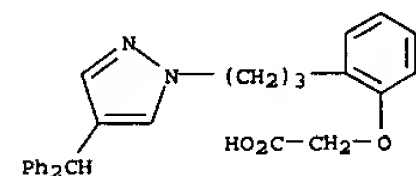


RN 152381-37-8 CAPLUS
 CN Acetic acid, [3-[2-[4-(diphenylmethyl)-1H-pyrazol-1-yl]ethyl]phenoxy]-(9CI) (CA INDEX NAME)

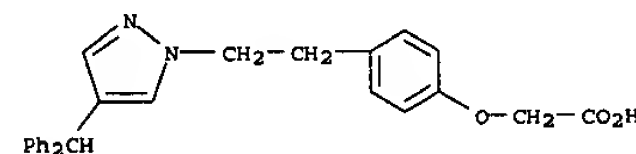
L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



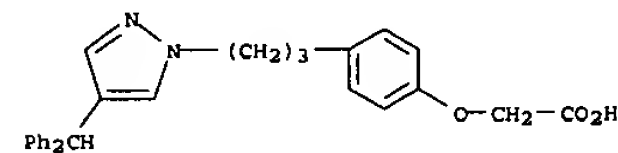
RN 152381-38-9 CAPLUS
 CN Acetic acid, [2-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-(9CI) (CA INDEX NAME)



RN 152381-40-3 CAPLUS
 CN Acetic acid, [4-[2-[4-(diphenylmethyl)-1H-pyrazol-1-yl]ethyl]phenoxy]-(9CI) (CA INDEX NAME)

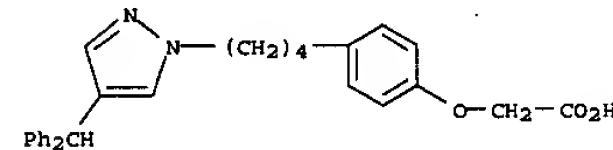


RN 152381-41-4 CAPLUS
 CN Acetic acid, [4-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-(9CI) (CA INDEX NAME)

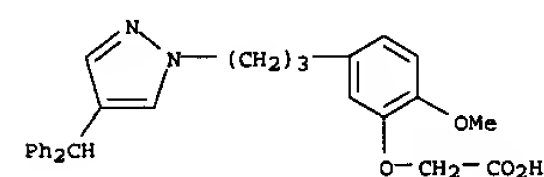


RN 152381-42-5 CAPLUS
 CN Acetic acid, [4-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-(9CI) (CA INDEX NAME)

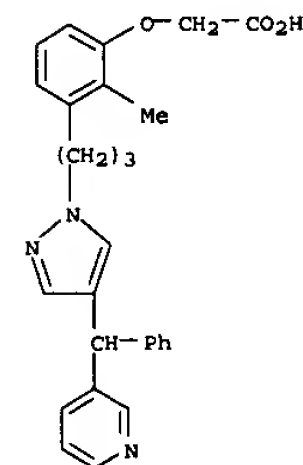
L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



RN 152381-46-9 CAPLUS
 CN Acetic acid, [5-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]-2-methoxyphenoxy]-(9CI) (CA INDEX NAME)



RN 153183-95-0 CAPLUS
 CN Acetic acid, [2-methyl-3-[3-[4-(phenyl-3-pyridinylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-(9CI) (CA INDEX NAME)

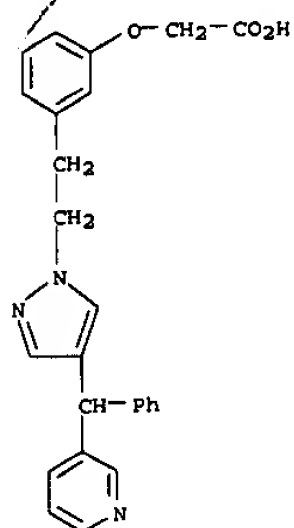


RN 153183-96-1 CAPLUS
 CN Acetic acid, [3-[2-[4-(phenyl-3-pyridinylmethyl)-1H-pyrazol-1-yl]ethyl]phenoxy]-(9CI) (CA INDEX NAME)

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L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)

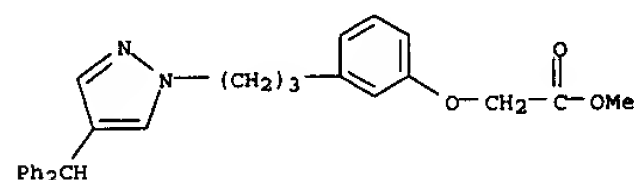


IT 152381-29-8P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. and antithrombotic and antihypertensive activities of,
reaction of)

RN 152381-29-8 CAPLUS

CN Acetic acid, [3-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-,
methyl ester (9CI) (CA INDEX NAME)



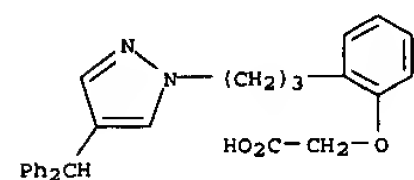
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152381-37-8P 152381-38-9P 152381-40-3P
152381-41-4P 152381-42-5P 152381-44-7P
152381-46-9P

RL: SPN (Synthetic preparation); PREP (Preparation)
(prepn. and antithrombotic and antihypertensive activity of)

RN 152381-30-1 CAPLUS

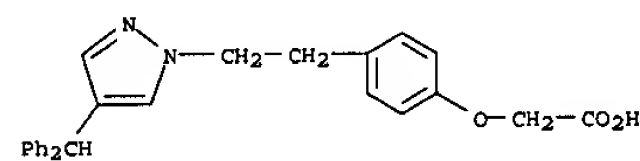
CN Acetic acid, [3-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-
(9CI) (CA INDEX NAME)

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



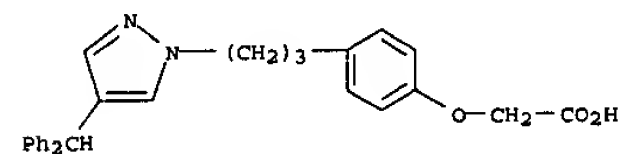
RN 152381-40-3 CAPLUS

CN Acetic acid, [4-[2-[4-(diphenylmethyl)-1H-pyrazol-1-yl]ethyl]phenoxy]-
(9CI) (CA INDEX NAME)



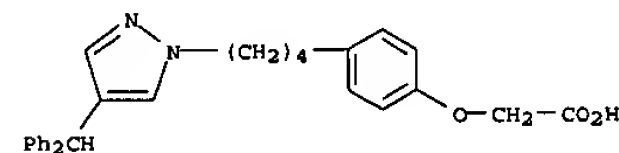
RN 152381-41-4 CAPLUS

CN Acetic acid, [4-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-
(9CI) (CA INDEX NAME)



RN 152381-42-5 CAPLUS

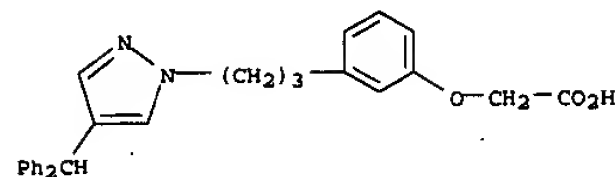
CN Acetic acid, [4-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-
(9CI) (CA INDEX NAME)



RN 152381-44-7 CAPLUS

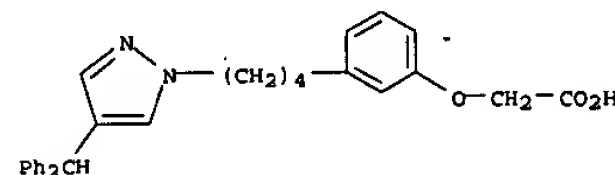
CN Acetic acid, [2-[4-[4-(phenyl-3-pyridinylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-
(9CI) (CA INDEX NAME)

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



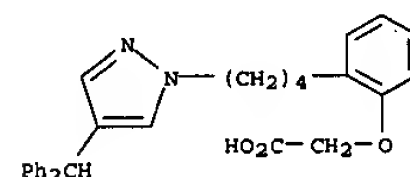
RN 152381-31-2 CAPLUS

CN Acetic acid, [3-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-
(9CI) (CA INDEX NAME)



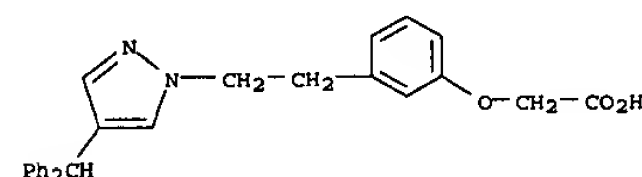
RN 152381-35-6 CAPLUS

CN Acetic acid, [2-[4-[4-(diphenylmethyl)-1H-pyrazol-1-yl]butyl]phenoxy]-
(9CI) (CA INDEX NAME)



RN 152381-37-8 CAPLUS

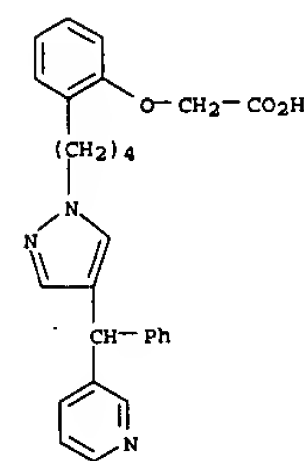
CN Acetic acid, [3-[2-[4-(diphenylmethyl)-1H-pyrazol-1-yl]ethyl]phenoxy]-
(9CI) (CA INDEX NAME)



RN 152381-38-9 CAPLUS

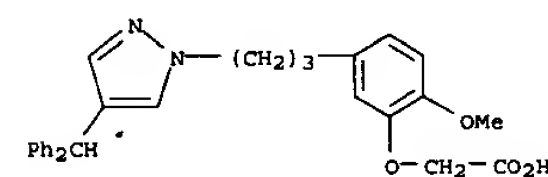
CN Acetic acid, [2-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]phenoxy]-
(9CI) (CA INDEX NAME)

L5 ANSWER 1 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



RN 152381-46-9 CAPLUS

CN Acetic acid, [5-[3-[4-(diphenylmethyl)-1H-pyrazol-1-yl]propyl]-2-
methoxyphenoxy]- (9CI) (CA INDEX NAME)



1000562

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1994:134381 CAPLUS

DOCUMENT NUMBER: 120:134381

TITLE: Nonpeptide angiotensin II antagonists derived from 1H-pyrazole-5-carboxylates and 4-aryl-1H-imidazole-5-carboxylates

AUTHOR(S): Ashton, Wallace T.; Hutchins, Steven M.; Greenlee, William J.; Doss, George A.; Chang, Raymond S. L.; Lotti, Victor J.; Faust, Kristie A.; Chen, Tsing Bau; Zingaro, Gloria J.; et al.

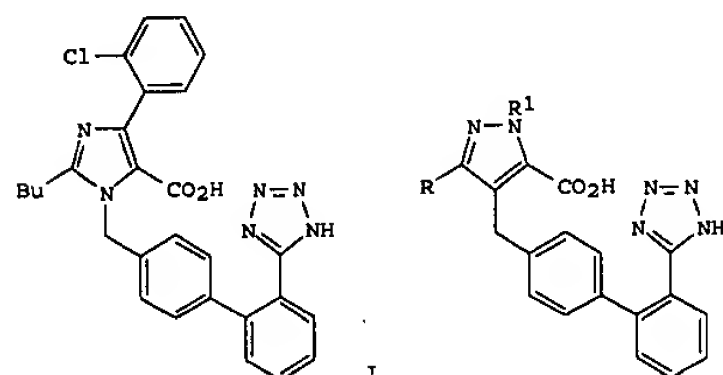
CORPORATE SOURCE: Merck Res. Lab., Rahway, NJ, 07065, USA

SOURCE: J. Med. Chem. (1993), 36(23), 3595-605

DOCUMENT TYPE: CODEN: JMCMAR; ISSN: 0022-2623

LANGUAGE: Journal

GI: English



AB Two series of potential angiotensin II antagonists derived from carboxyl-functionalized "diazole" heterocycles have been prepd. and evaluated. Initially, a limited investigation of 4-arylimidazole-5-carboxylates led to 2-n-butyl-4-(2-chlorophenyl)-1-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-imidazole-5-carboxylic acid (I), which was found to be a highly potent antagonist of the rabbit aorta AT1 receptor (IC50 0.55 nM). In conscious, normotensive rats, I at 0.1 mg/kg i.v. inhibited the pressor response to AII by 88%, with a duration of >6 h. More extensively studied was an isosteric series of 3-alkyl-4-[[2'-(1H-tetrazol-5-yl)biphenyl-4-yl]methyl]-1H-pyrazole-5-carboxylates bearing aryl, alkyl, or aralkyl substituents at N1. These compds. were available in highly regioselective fashion via condensation of a substituted hydrazine hydrochloride with a 2-(methoxyimino)-4-oxoalkanoate intermediate. In vitro, the most potent pyrazolecarboxylic acids were II (R = Bu; R1 = 2,6-dichlorophenyl, 2-(trifluoromethyl)phenyl, benzyl, and phenethyl), all with IC50 values of 0.18-0.24 nM. Although less potent

in the receptor assay, 3-n-propylpyrazolecarboxylic acids were at least as effective as their Bu counterpart in vivo. Several of the pyrazolecarboxylic acid derivs. demonstrated potent, long-lasting oral activity in rats. At 1 mg/kg po, the II (R = Bu, R1 = benzyl; R = Pr, R1

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)

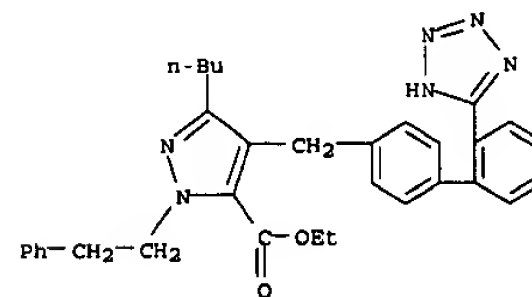
= 2,6-dichlorophenyl, 2,2,2-trifluoroethyl, and benzyl) analogs all gave >75% inhibition of the AII pressor response in the rat model, with duration of action >23 h.

IT 152713-37-6P 152713-50-3P

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. and angiotensin II antagonist activity of)

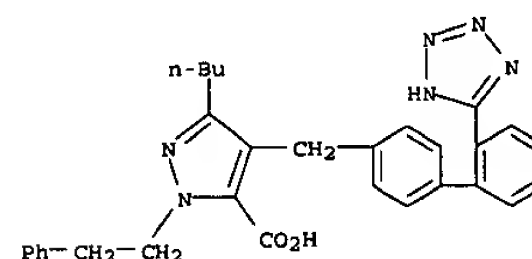
RN 152713-37-6 CAPLUS

CN 1H-Pyrazole-5-carboxylic acid, 3-butyl-1-(2-phenylethyl)-4-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]-, ethyl ester (9CI) (CA INDEX NAME)



RN 152713-50-3 CAPLUS

CN 1H-Pyrazole-5-carboxylic acid, 3-butyl-1-(2-phenylethyl)-4-[[2'-(1H-tetrazol-5-yl)[1,1'-biphenyl]-4-yl]methyl]- (9CI) (CA INDEX NAME)



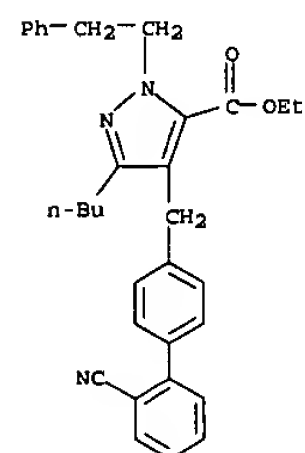
IT 152713-71-8P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation) (prepn. and cyclization of, with azide, triazole deriv. from)

RN 152713-71-8 CAPLUS

CN 1H-Pyrazole-5-carboxylic acid, 3-butyl-4-[[2'-(cyano[1,1'-biphenyl]-4-yl)methyl]-1-(2-phenylethyl)-, ethyl ester (9CI) (CA INDEX NAME)

L5 ANSWER 2 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)



L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS

ACCESSION NUMBER: 1993:603377 CAPLUS

DOCUMENT NUMBER: 119:203377

TITLE: reaction of N-substituted acetohydrazides with 2-substituted cinnamitriles. Competitive cyclizations to pyrazolo[3,4-b]pyridinones and [1,2,4]triazolo[1,5-a]pyridinones

AUTHOR(S): Hadi, Ali; Martin, Nazario; Seoane, Carlos; Soto, Jose

CORPORATE SOURCE: Fac. Quim., Univ. Complutense, Madrid, 28040, Spain

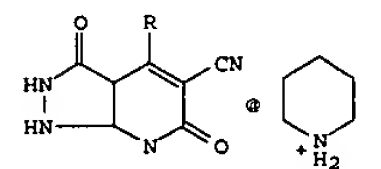
SOURCE: J. Chem. Soc., Perkin Trans. 1 (1993), (9), 1045-50

DOCUMENT TYPE: CODEN: JCPRB4; ISSN: 0300-922X

LANGUAGE: Journal

OTHER SOURCE(S): English

GI: CASREACT 119:203377



AB A novel prepn. of pyrazolo[3,4-b]pyridinones I (R = aryl) from 2'-acyl-2-cyanoacetohydrazide and arylidenecyanoacetates is described.

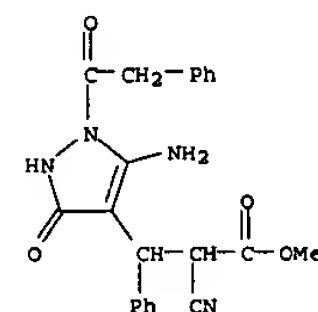
In the reaction, an alternative cyclization, leading to [1,2,4]triazolo[1,5-a]pyridinones takes place. Compds. I were isolated from the reaction mixt. as the corresponding.

IT 150568-54-0P 150568-55-1P 150568-57-3P

RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)

RN 150568-54-0 CAPLUS

CN 1H-Pyrazole-4-propanoic acid, 5-amino-.alpha.-cyano-2,3-dihydro-3-oxo-.beta.-phenyl-1-(phenylacetyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 150568-55-1 CAPLUS

CN 1H-Pyrazole-4-propanoic acid, 5-amino-.alpha.-cyano-2,3-dihydro-3-oxo-.beta.-phenyl-1-(phenylacetyl)-, methyl ester, compd. with piperidine

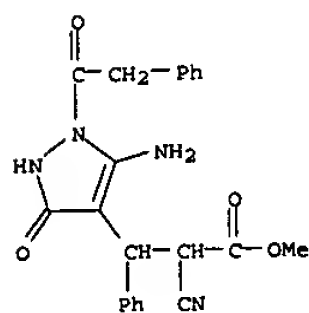
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L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)
(1:1) (9CI) (CA INDEX NAME)

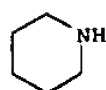
CM 1

CRN 150568-54-0
CMP C22 H20 N4 O4



CM 2

CRN 110-89-4
CMP C5 H11 N



RN 150568-57-3 CAPLUS
CN 1H-Pyrazole-4-propanoic acid, 5-amino-.alpha.-cyano-2,3-dihydro-3-oxo-.beta.-phenyl-1-(phenylacetyl)-, ethyl ester, compd. with piperidine

(1:1)
(9CI) (CA INDEX NAME)

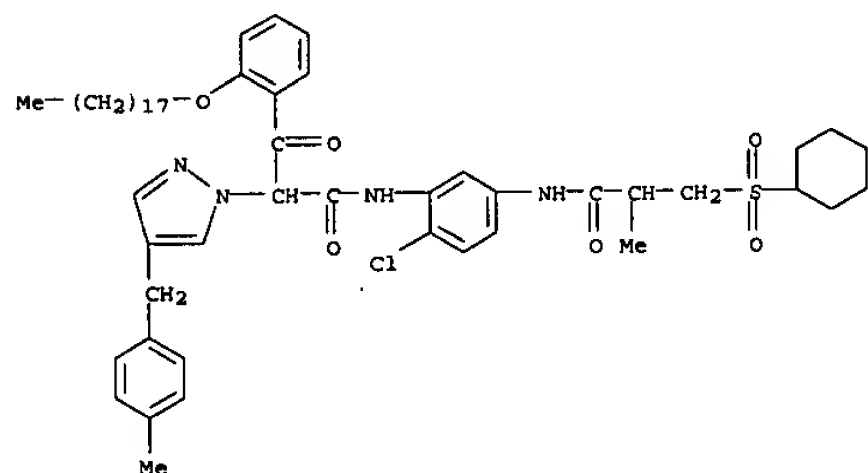
CM 1

CRN 150568-56-2
CMP C23 H22 N4 O4

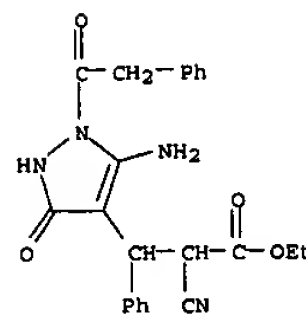
L5 ANSWER 4 OF 5 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER: 1988:539046 CAPLUS
DOCUMENT NUMBER: 109:139046
TITLE: Silver halide photographic material containing yellow coupler
INVENTOR(S): Tsuruta, Mayumi; Mizukura, Noboru; Nakagawa, Satoshi
PATENT ASSIGNEE(S): Konica Co., Japan
SOURCE: Jpn. Kokai Tokkyo Koho, 16 pp.
CODEN: JKXXAF
DOCUMENT TYPE: Patent
LANGUAGE: Japanese
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 63092951	A2	19880423	JP 1986-238222	19861007

GI For diagram(s), see printed CA Issue.
AB In the title photog. material, .gtoreq.1 of photog. Ag halide emulsion layers contains a yellow coupler I [R1 = alkyl, cycloalkyl, aryl; R2 = group which can be substituted to the benzene ring; R3 = H, alkyl, aryl, heterocyclyl; X = alkylene, cycloalkylene, arylene, alkylene arylene, arylene alkylene, or -A-V-B- (A, B = alkylene, arylene, alkylenearylene, or arylmealkylene; V = divalent connecting group); Y = alkyl, cycloalkyl, aryl, heterocyclyl; Z = nonmetal atoms to form a 5- or 6-membered ring with -N(CO)n-; m = 0, 1; n = 0-2]. The photog. material shows improved color-forming d., reduced fog, and improved storage stability.
IT 116624-91-0
RL: TEM (Technical or engineered material use); USES (Uses) (photog. yellow coupler)
RN 116624-91-0 CAPLUS
CN 1H-Pyrazole-1-acetamide, N-[2-chloro-5-[[3-(cyclohexylsulfonyl)-2-methyl-1-oxopropyl]amino]phenyl]-4-[(4-methylphenyl)methyl]-.alpha.-[2-(octadecyloxy)benzoyl]- (9CI) (CA INDEX NAME)

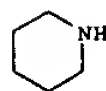


L5 ANSWER 3 OF 5 CAPLUS COPYRIGHT 2002 ACS (Continued)

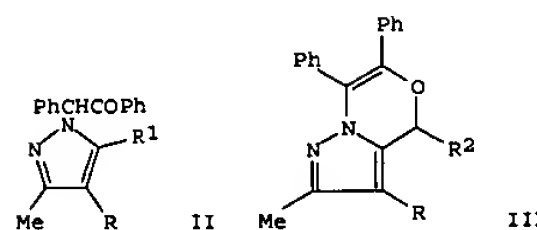


CM 2

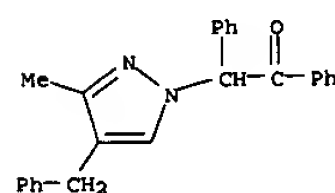
CRN 110-89-4
CMP C5 H11 N



L5 ANSWER 5 OF 5 CAPLUS COPYRIGHT 2002 ACS
ACCESSION NUMBER: 1984:423412 CAPLUS
DOCUMENT NUMBER: 101:23412
TITLE: Reactions of azines. 8. Synthesis and thermal rearrangement of 1-oxo-3,4-diaza-2,4,6-heptatrienes and 1-oxo-3,4-diaza-2,4,6,7-octatetraenes (allenyl azines)
AUTHOR(S): Schweizer, Edward E.; Lee, Kee Jung
CORPORATE SOURCE: Dep. Chem., Univ. Delaware, Newark, DE, 19711, USA
SOURCE: J. Org. Chem. (1984), 49(11), 1959-64
CODEN: JOCEAH; ISSN: 0022-3263
DOCUMENT TYPE: Journal
LANGUAGE: English
OTHER SOURCE(S): CASREACT 101:23412
GI



AB Cycloaddn. reactions of 1-oxo-3,4-diaza-2,4,6-heptatrienes, obtained from PhCOCPh:NN:CMcCR:PPh3 (I; R = Me, Et, Pr, H2C:CHCH2, PhCH2), with aldehydes gave substituted pyrazoles II (R = Me, R1 = Ph, p-O2NC6H4; R = Et, Pr, R1 = p-O2NC6H4; R = H2C:CHCH2 R1 = Ph; R = PhCH2, R1 = H) in 66-89% yield. I (R = PhCO) failed in the olefination reaction, giving only the corresponding acetylene. A similar allenylation reaction of I (R = Me, H2C:CHCH2, PhCH2) with R2HC:CO (R2 = H, Ph, PhCH2) gave pyrazolo[5,1-c][1,4]oxazines III in 65-81% yield via the intermediate 1-oxo-3,4-diaza-2,4,6,7-octatetraenes. I (R = PhCO) only reacted with H2C:CO to give III in 21% yield.
IT 89849-25-2P
RL: SPN (Synthetic preparation); PREP (Preparation) (prepn. of)
RN 89849-25-2 CAPLUS
CN Ethanone, 2-[3-methyl-4-(phenylmethyl)-1H-pyrazol-1-yl]-1,2-diphenyl- (9CI) (CA INDEX NAME)



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=> file reg

COST IN U.S. DOLLARS

SINCE FILE

TOTAL

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SESSION

FULL ESTIMATED COST

23.93

164.42

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)

SINCE FILE

TOTAL

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SESSION

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STRUCTURE FILE UPDATES: 7 MAY 2002 HIGHEST RN 412267-09-5

DICTIONARY FILE UPDATES: 7 MAY 2002 HIGHEST RN 412267-09-5

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Calculated physical property data is now available. See HELP PROPERTIES
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Registry File, for complete details:

<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>Testing the current file.... screen

ENTER SCREEN EXPRESSION OR (END):end

=>

Uploading C:\Program Files\Stnexp\Queries\0989932211.str

L6 STRUCTURE UPLOADED

=> que L6

L7 QUE L6

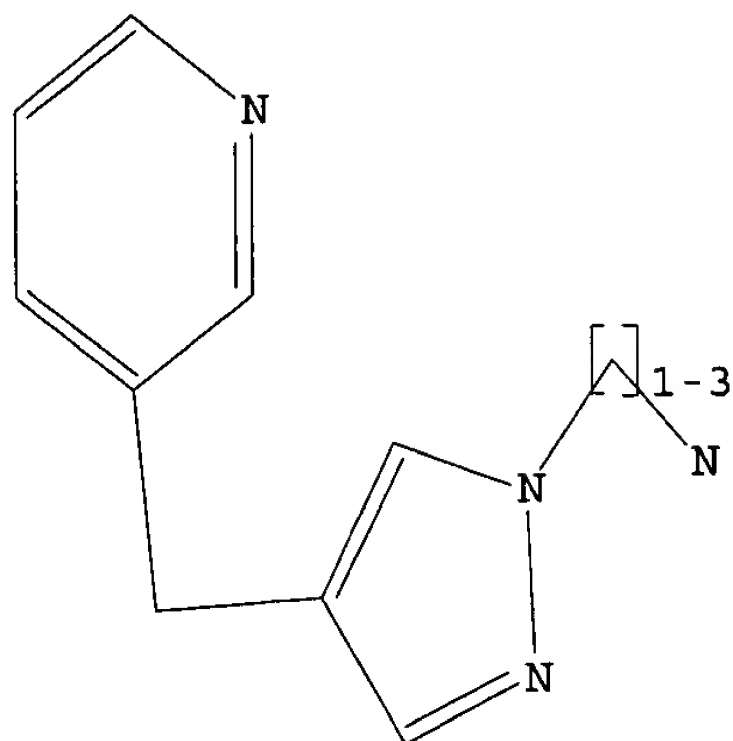
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L7 HAS NO ANSWERS

L6 STR

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Structure attributes must be viewed using STN Express query preparation.
L7 QUE ABB=ON PLU=ON L6

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SAMPLE SEARCH INITIATED 12:32:10 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 0 TO ITERATE

100.0% PROCESSED 0 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 0 TO 0
PROJECTED ANSWERS: 0 TO 0

L8 0 SEA SSS SAM L6

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FULL SEARCH INITIATED 12:33:08 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 101 TO ITERATE

100.0% PROCESSED 101 ITERATIONS 22 ANSWERS
SEARCH TIME: 00.00.01

L9 22 SEA SSS FUL L1

=> s 17 full
FULL SEARCH INITIATED 12:33:31 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 8 TO ITERATE

100.0% PROCESSED 8 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.02

L10 0 SEA SSS FUL L6

=> logoff
ALL L# QUERIES AND ANSWER SETS ARE DELETED AT LOGOFF
LOGOFF? (Y)/N/HOLD:y

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COST IN U.S. DOLLARS	SINCE FILE	TOTAL
	ENTRY	SESSION
FULL ESTIMATED COST	280.94	445.36
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE	TOTAL
	ENTRY	SESSION
CA SUBSCRIBER PRICE	0.00	-3.10

STN INTERNATIONAL LOGOFF AT 12:33:49 ON 09 MAY 2002

Kamal Saeed